Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application.

- 1. Cancelled.
- 2. (Currently Amended) A non-tumorigenic cell composition derived from embryonic stem cells, the composition comprising about 100% isolated <u>neural cells and</u> neural precursor cells, the neural precursor cells having the ability to differentiate into neuronal cells[[,]] <u>or</u> glial cells, <u>or combinations thereof</u>,

the composition being obtainable by:

- (a) culturing the embryonic stem cells to produce neural precursor cells;
- (b) culturing the neural precursor cells from (a) in a first growth factorcontaining serum-free medium;
- (c) culturing the cells from (b) in a second growth factor-containing serumfree medium; and
- (d) culturing the cells from (c) in a third growth factor-containing serum-free medium,

wherein the cultured cells from (d) are non-tumorigenic and <u>comprise neural</u> <u>cells and neural precursor cells</u>, <u>wherein the neural precursor cells</u> have the ability to differentiate into neuronal cells[[,]] <u>or</u> glial cells, <u>or combinations thereof</u>.

- 3. (Previously Presented) The cell composition according to claim 2, wherein the embryonic stem cells in step (a) are in the form of cell aggregates.
- 4. Cancelled.
- 5. Cancelled.

- 6. (Previously Presented) The cell composition according to claim 2, wherein the cells of steps (c) and (d) grow as a monolayer.
- 7. Cancelled.
- 8. (Currently Amended) The cell composition according to claim 2, comprising cells with neuronal, astroglial or oligodendroglial properties, or a combination thereof.
- 9. (Previously Presented) The cell composition according to claim 2, wherein the embryonic stem cells are obtained after nuclear transfer into oocytes.
- 10. (Previously Presented) The cell composition according to claim 2, wherein the embryonic stem cells are obtained from embryonic germ cells.
- 11. (Previously Presented) The cell composition according to claim 2, wherein the cells are mammalian cells.
- 12. (Previously Presented) The cell composition according to claim 11, wherein the cells are isolated from a mammal selected from the group consisting of mouse, rat, hamster, pig, cow, primate, and human.
- 13. Cancelled.
- 14. Cancelled.
- 15. (Previously Presented) A cell library comprising autologous and non-autologous cells according to claim 47.
- 16. 45. Cancelled.
- 46. (Previously Presented) A pharmaceutical composition comprising the precursor cells of claim 47.
- 47. (Currently Amended) A non-tumorigenic cell composition derived from embryonic stem cells,

the composition comprising about 100% isolated <u>neural cells and</u> neural precursor cells, the neural precursor cells having the ability to differentiate into neuronal cells[[,]] <u>or</u> glial cells, <u>or combinations thereof</u>, and wherein the cell composition is non-tumorigenic.

- 48. (Previously Presented) The cell composition of claim 2, wherein the embryonic stem cells in (a) are cultured in serum-free medium.
- 49. Cancelled.
- 50. (Previously Presented) The cell composition of claim 3, wherein the cell aggregates are embryoid bodies.
- 51. Cancelled.
- 52.-75. Not entered.
- 76. (Currently Amended) A non-tumorigenic cell composition derived from embryonic stem cells, the composition comprising about 100% isolated <u>neural cells and</u> neural precursor cells that have the ability to differentiate into neuronal cells[[,]] <u>or</u> glial cells, <u>or combinations thereof</u>, the composition being obtainable by:
 - (a) culturing the embryonic stem cells to produce neural precursor cells;
 - (b) culturing the neural precursor cells from (a) in a first growth factorcontaining serum-free medium; and
 - (c) culturing the cells from (b) in a second growth factor-containing serumfree medium to produce neural spheres,

wherein the cells of the neural spheres are non-tumorigenic and <u>comprise neural</u> <u>cells and neural precursor cells</u>, <u>wherein the neural precursor cells</u> have the ability to differentiate into neuronal cells, astrogilal cells, <u>or</u> oligodendroglial cells, or <u>combinations thereof</u>.

- 77. (Previously Presented) The cell composition according to claim 76, wherein the embryonic stem cells in (a) are in the form of cell aggregates.
- 78. (Previously Presented) The cell composition of claim 77, wherein the cell aggregates are embryoid bodies.
- 79. (Previously Presented) The cell composition of claim 76, wherein the embryonic stem cells in (a) are cultured in serum-free medium.
- 80. (Previously Presented) The cell composition according to claim 76, wherein the embryonic stem cells are obtained after nuclear transfer into oocytes.
- 81. (Previously Presented) The cell composition according to claim 76, wherein the embryonic stem cells are obtained from embryonic germ cells.
- 82. (Previously Presented) The cell composition according to claim 76, wherein the cells are mammalian cells.
- 83. (Previously Presented) The cell composition according to claim 82, wherein the cells are isolated from a mammal selected from the group consisting of mouse, rat, hamster, pig, cow, primate, and human.
- 84. Cancelled.
- 85. (Previously Presented) A cell library comprising cells according to claim 76, which are autologous and nonautologous cells.
- 86. (Previously Presented) A pharmaceutical composition comprising the precursor cells of claim 76.
- 87. (Currently Amended) A non-tumorigenic cell composition derived from embryonic stem cells, the composition comprising about 100% isolated <u>neural cells and</u> neural precursor cells that have the ability to differentiate into glial cells, the composition being obtainable by:

- (a) culturing the embryonic stem cells to produce neural precursor cells;
- (b) culturing the neural precursor cells from (a) in a first growth factorcontaining serum-free medium;
- (c) culturing the cells from (b) in a second growth factor-containing serum-free medium to produce neural spheres; and
- (d) culturing the neural spheres from (c) in a third growth factorcontaining serum-free medium to produce a monolayer of glial precursor cells,

wherein the cells of the monolayer are non-tumorigenic and <u>comprise neural</u> <u>cells and neural precursor cells, wherein the neural precursor cells</u> have the ability to differentiate into glial cells.

- 88. (Previously Presented) The cell composition according to claim 87, wherein the embryonic stem cells in step (a) are in the form of cell aggregates.
- 89. (Previously Presented) The cell composition of claim 88, wherein the cell aggregates are embryoid bodies.
- 90. (Previously Presented) The cell composition of claim 87, wherein the embryonic stem cells in (a) are cultured in serum-free medium.
- 91. (Previously Presented) The cell composition according to claim 87, wherein the embryonic stem cells are obtained after nuclear transfer into oocytes.
- 92. (Previously Presented) The cell composition according to claim 87, wherein the embryonic stem cells are obtained from embryonic germ cells.
- 93. (Previously Presented) The cell composition according to claim 87, wherein the cells are mammalian cells.

- 94. (Previously Presented) The cell composition according to claim 93, wherein the cells are isolated from a mammal selected from the group consisting of mouse, rat, hamster, pig, cow, primate, and human.
- 95. Cancelled.
- 96. (Previously Presented) A cell library comprising cells according to claim 87, which are autologous and nonautologous cells.
- 97. (Previously Presented) A pharmaceutical composition comprising the precursor cells of claim 87.
- 98. (Previously Presented) A cell library comprising cells according to claim 2, which are autologous and nonautologous cells.
- 99. (Previously Presented) A pharmaceutical composition comprising the precursor cells of claim 2.
- 100. (New) The cell composition according to claim 2, wherein the second growth factor-containing serum-free medium comprises bFGF and EGF.
- 101. (New) The cell composition according to claim 2, wherein the third growth factor-containing serum-free medium comprises bFGF and PDGF.
- 102. (New) The cell composition according to claim 76, wherein the second growth factor-containing serum-free medium comprises bFGF and EGF.
- 103. (New) The cell composition according to claim 87, wherein the second growth factor-containing serum-free medium comprises bFGF and EGF.
- 104. (New) The cell composition according to claim 87, wherein the third growth factor-containing serum-free medium comprises bFGF, EGF, or a combination thereof.